

BERSHADSKIT, A.L., doktor tekhnicheskikh nauk; MDISMIN, A.V.

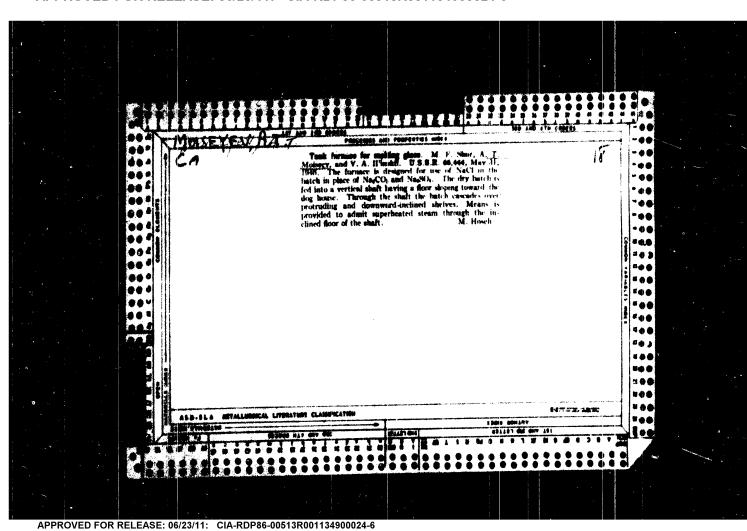
Regulating frame saw tension. Der. prom. 6 no.316-8 Mr 57, (MLRA 10:5)

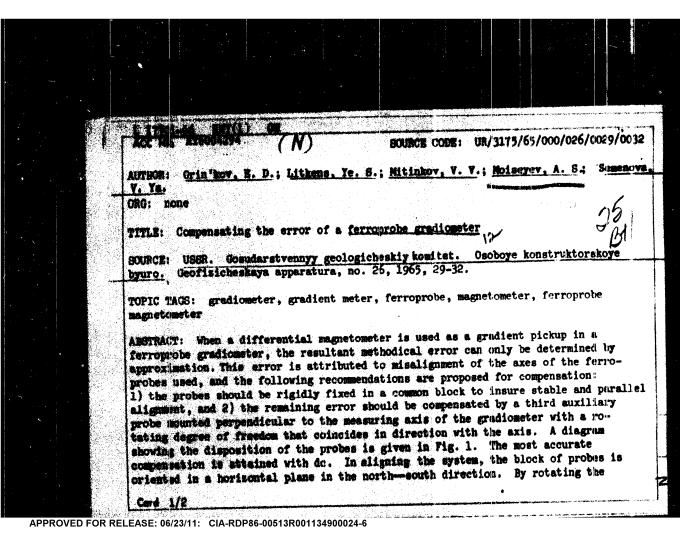
1. Belorusskiy lesotekhnicheskiy institut im. S.M. Kirova. (Band saws)

MCCOTYSM. A. T., Eng.
Wells
New wall block constructions. Gcr. khoz.Mosk. 26 No. 7, 1952

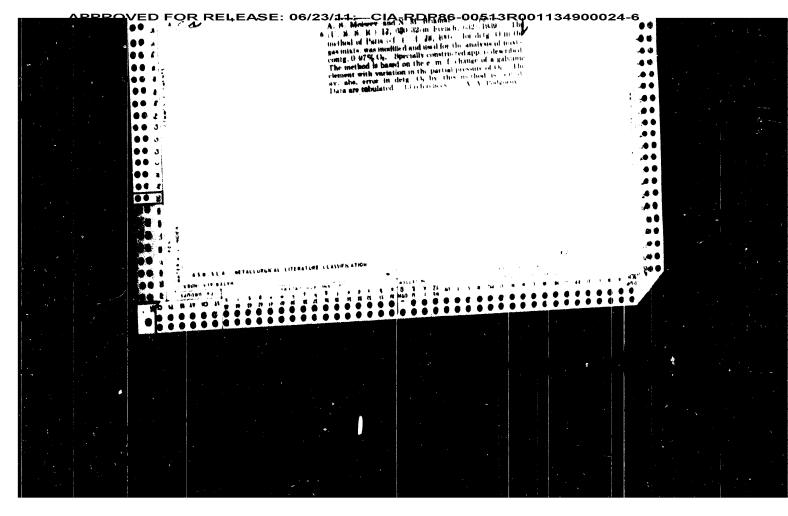
Monthly List of Russian Accessions, Library of Congress, December 1967. Unclassified

APPROVED FOR RELEASE: 06/23/11: CIA-RDP86-00513R001134900024-6

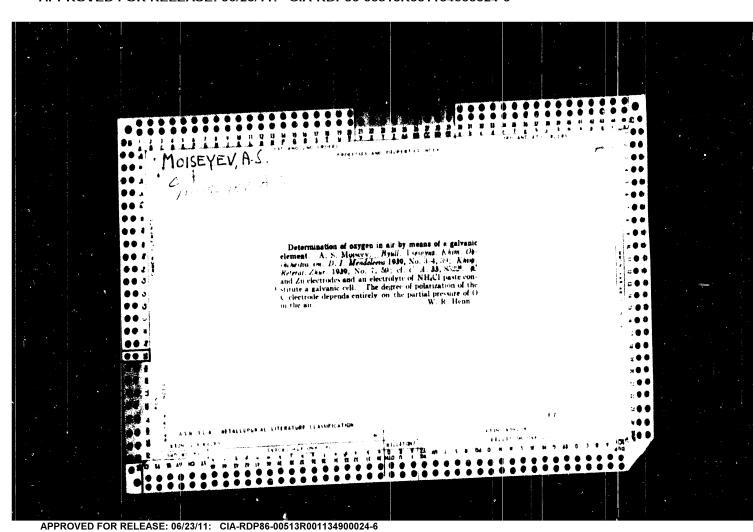




ROVED FOR RELEASE: 06/23/11: CIA-RDP86-00513R001134900024-6 Jennyral Vol'f, A. S. Moiseyev, P. V. Korystin, and I. V. Novik, Vodosnabzheniye i Sanitarnaya Tekhnika, No 12, Dec 56, pp 8-10 The article gives a brief history of the development of portable ionite filters for purification (elimination of salts and impurities) from water to render it potable, conducted by the All-Union Scientific Mesearch Institute for Hydraulic Engineering and Sanitary Engineering Works, from 1950 to present. The article also describes in detail the construction and character. istics of a portable ionite water filter developed in 1955 by the above institute in conjunction with the Novosibirsk Scientific Research Sanitary Institute, the filter being designed for the use of small groups under field conditions in areas of high mineral content. The purified water output of the filter on a single charge of ionites is 250 liters when the salt content of the original water is less than 3 g/l. When the original salt concentration is 5-6 g/l, the fresh water cutput is reduced to 100-120 1. The filtering unit itself is cylindrical in shape, the dimensions SUM. 1287 being one meter x 200 mm.



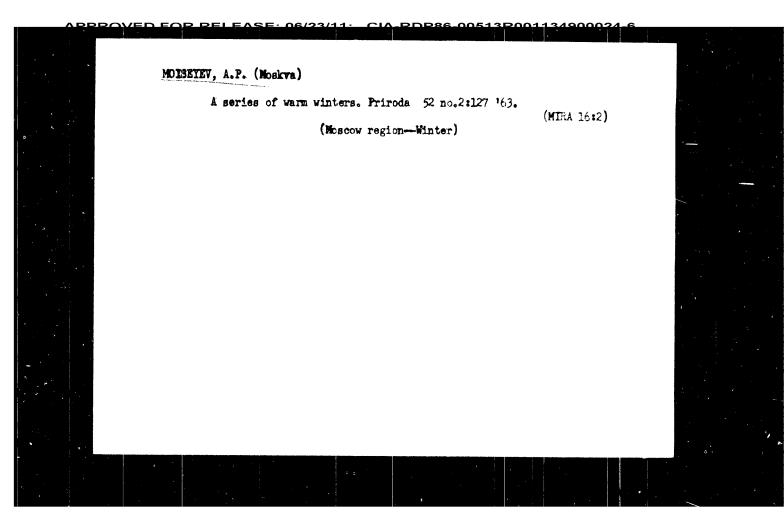
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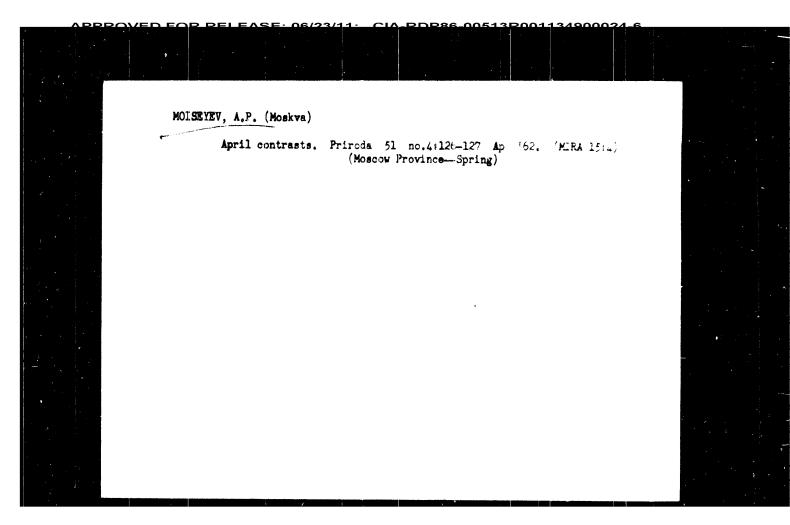


MOISHIW, Aleksandr Sergeyevich, inshemer; SHVETSOV, I.B., redaktor;

[Achievements of Soviet agricultural machinery construction under the fifth five-year plan] Dostishenia sovetskogo sel'spothostaist-vennogo mashinostroenia v plato! platiletko. Moskva, Isd-vo "Zmannie." 1954. 29 p. (Veseclusnoe obshchestvo po resprestrameniiu politicheskith in machaphra manii. Ser. 4, no. 19) (MERA 7:9)

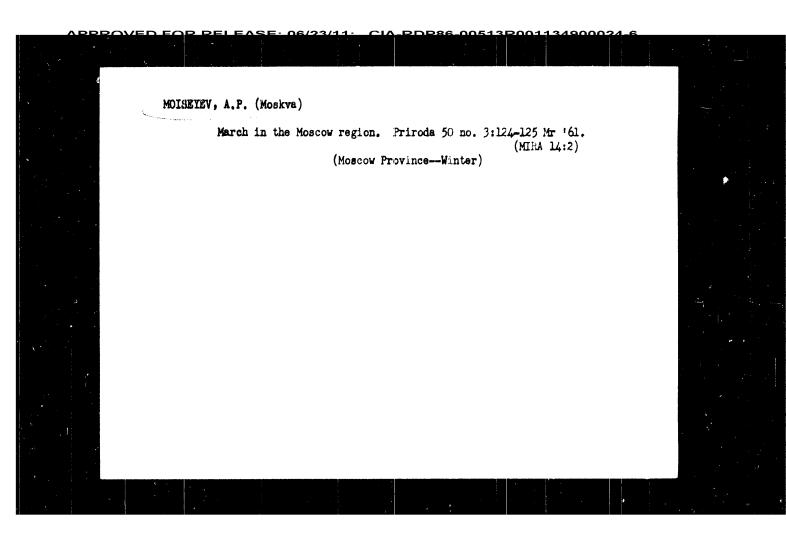
(Agricultural machinery industry)

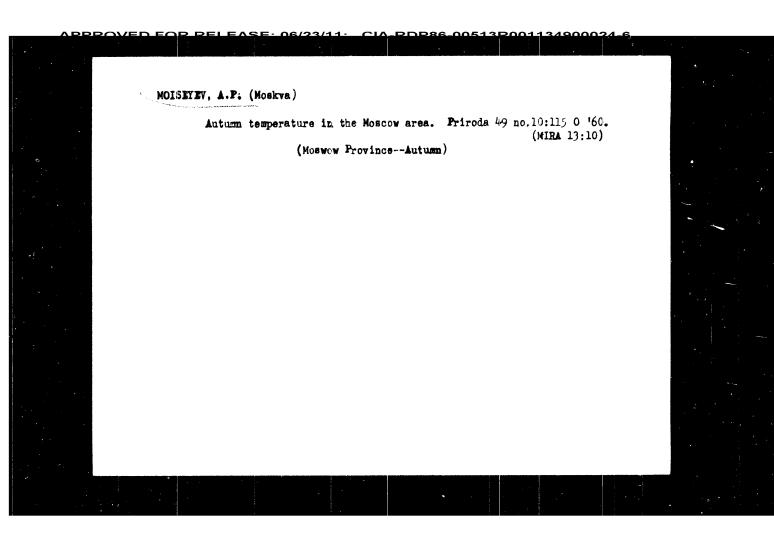




MOISEYEV, A.P.; KUVSHINOVA, K.V., kand. eograf.nauk December in Moscow. Priroda 50 no.12:123-124 D (MIRA 14:12) 1. Moskerskiy filial Geograficheskogo obshchestva SSSR (for Moiseyev). 2. Institut geografii AN SSSR, Moskva (for Kuvshinova).

(Moscow--Winter)





AUTHOR:

Moiseyev, A.P. (Moscow)

SOV-26-58-8-51/51

TITLE:

Migration of Martins to the Countryside of Moscow (Otlet

strizhey v Podmoskov'ye)

PERIODICAL:

Priroda, 1958, Nr 8, p 127 (USSR)

ABSTRACT:

Observations between 1934 and 1956 showed that martins are seen passing southward through the countryside of Moscow in the first half of August, 11 August being the mean date. On 13 August 1936 a flock of one thousand was noticed at a distance of 1 to 2 km in the village of Kolomenskoye of the Leninskiy District. The latest migration date of a major flock of martins was recorded on 6 Sep 1953. Individual martins were seen on 24 Sep 1957 and even on 3 Oct. These happened to be unusually warm months followed by a cold in the first decade of the October month that surpassed the conditions of the ensuing first tendays of November.

1. Martins--USSR

Card 1/1

USCOMM-DC-55419

AUTHOR: Moiseyev, A.P. (Moscow)

FITLE: A Summer of Thunderstorms and Tempests in the Moscow area (Leto groz i cur' v podmoskov'ye)

PERIODICAL: Priroda, 1958, Nr 7, p 126 (USSR)

ABSTRACT: In the summer of 1957 there were 36 days with thunderstorms, often accompanied by tempests and hall in the Moscow area. The mean annual number of thunderstorm days is 21.

1. Storms--USSR

Card 1/1

ACTRON: Moregov, 7:1 Correr)

TITLE: Cummer Full Yoons (Letsiye polablumiye)

PARIODICAL: Driroda, 1959, Nr 6, p 175-126 (USIR)

ASCTRACT: The article deals with the summer full moon and its visitility in the central left of the USI. It is pointed out that the moon does not rise very high above the horizon, which is two to certain negaliarities of the moon's circulation around the earth and their joint circulation around the sur.

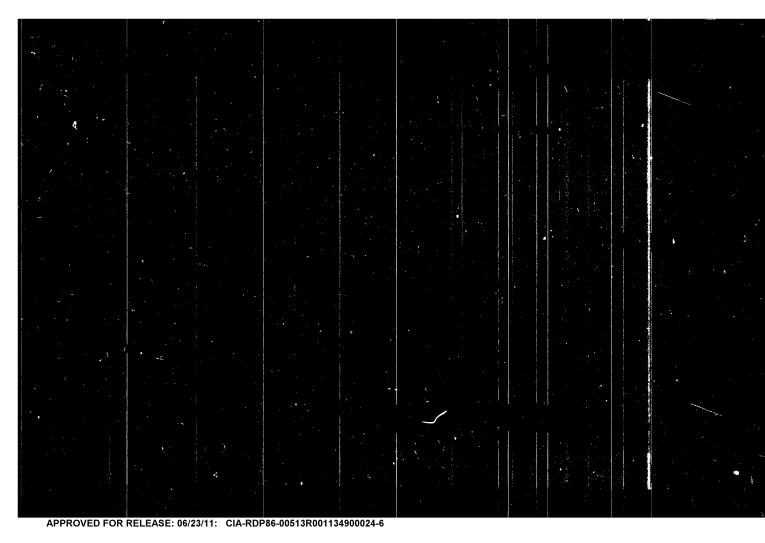
Card 1/1 There is one table and one graph.

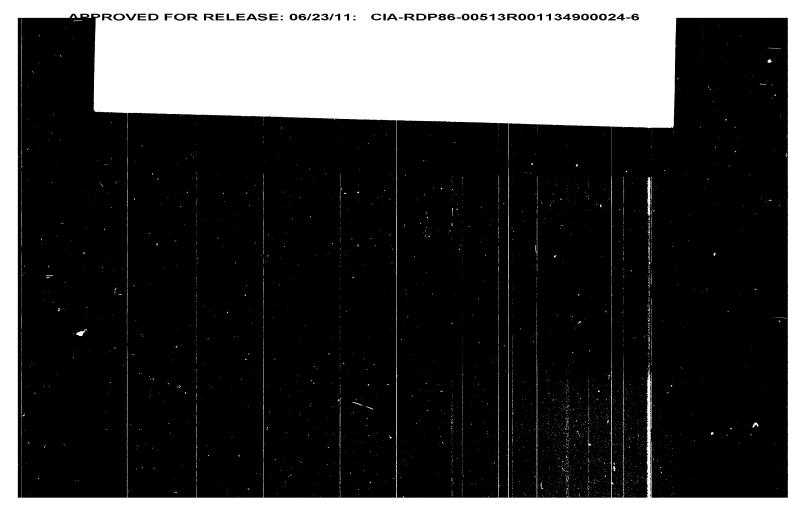
1. Moon-Visibility-USSR

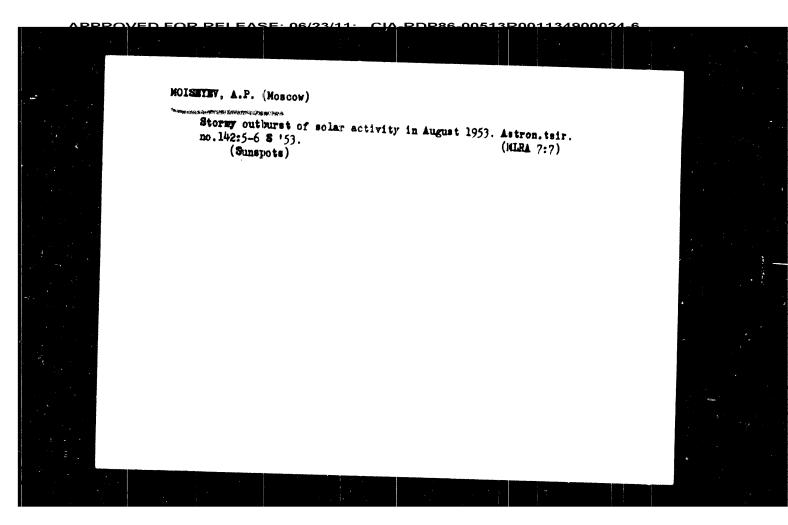
SOV-26-58-3-48/51 AUTHORS: Moiseyev, A.P. (Moscow), Kuznitsya, E.A. (Slobodskey, Kirov Oblast') TITLE: Early Spring Thunderstorms (Ranniye vesenniye grozy) Priroda, 1958, Nr 3, pp 125-126 (USSR) PERIODICAL: ABSTRACT: Thunderstorms during the cold season are comparatively rare in the European continental regions of the USSR. If they happen, it is due to the invasion of intensive cyclones with Atlantic air masses. This is accompanied by thawing, fogs and violent winds causing drifting snowstorms. 2. Thunderclouds--USSR 3. Cyclones--Meteoro-1. Storms--USSR logical effects Card 1/1

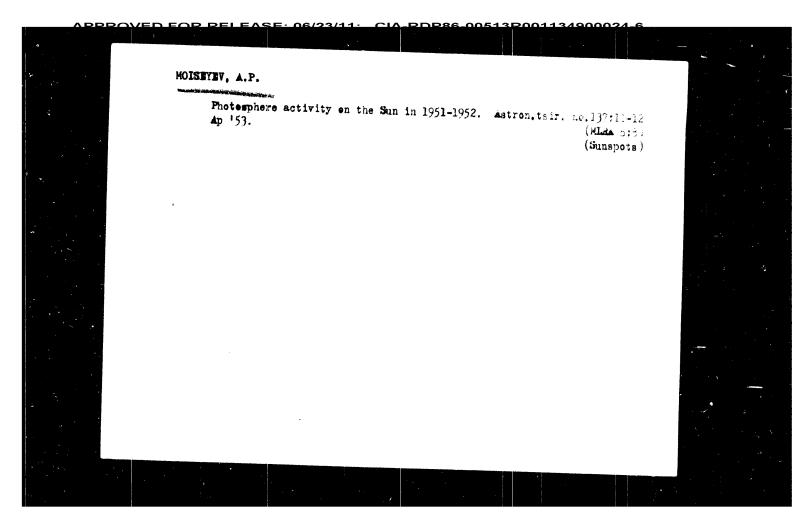
ME) ST-YEV P. 1. 16-08-1-40/18 AUTHOR: Galakhov, N.N., Doctor of Reographical Colenges: Moiseyev, 2000. (Moscow); Klintsov, A.F. (Bolinsk, Sakhalin oblast' TITLE: Calendar of Nature (Kalendar' prirody) PERIODICAL: Priroda, 1958, Nr 2, pp 126-128 (USSR) These three separate reports deal with the average and exception-ABSTRACT: al temperatures for February in the MSSR as a whole and around Moscow and on Sakhalin in particular. There is 1 diagram. ASSOCIATION: Institut geografii Akademii nauk BSSR (Institute of Reography of the USSR Academy of Sciences, Moscow) Card 1/11. Meteorology--USSR 2. Temperature--Applications

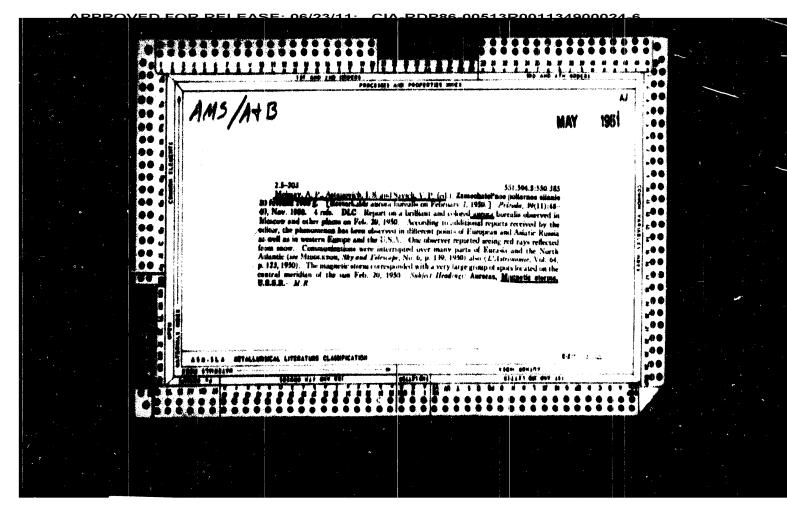
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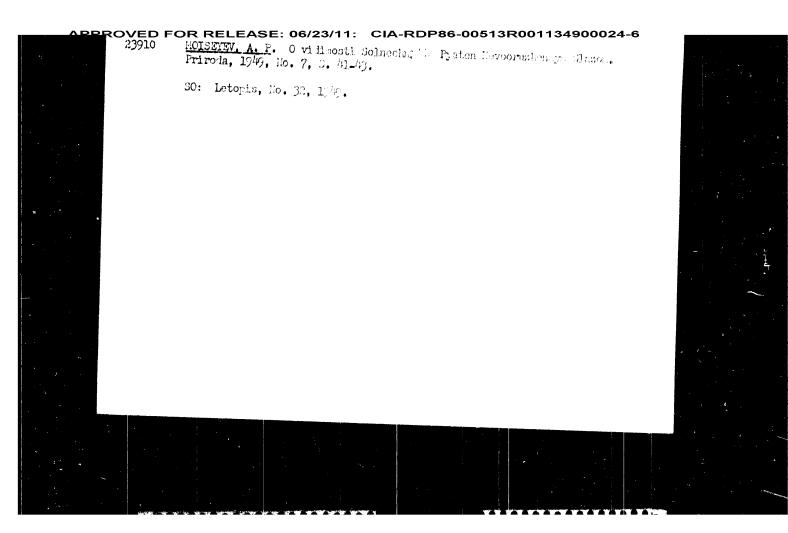


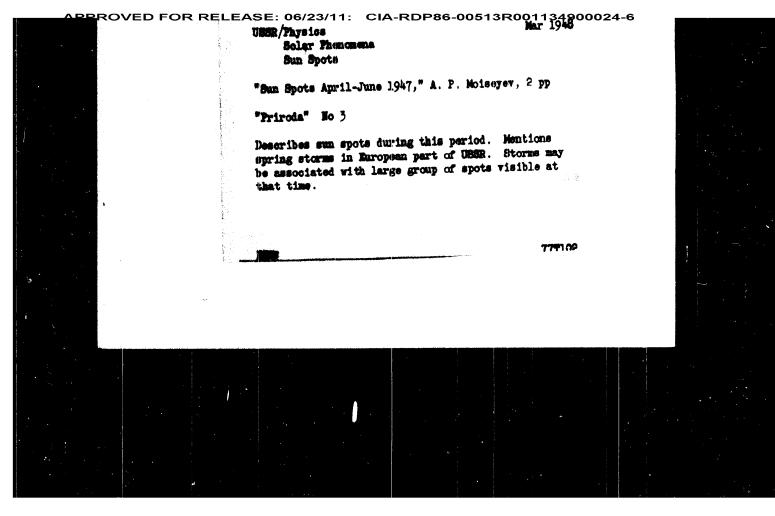


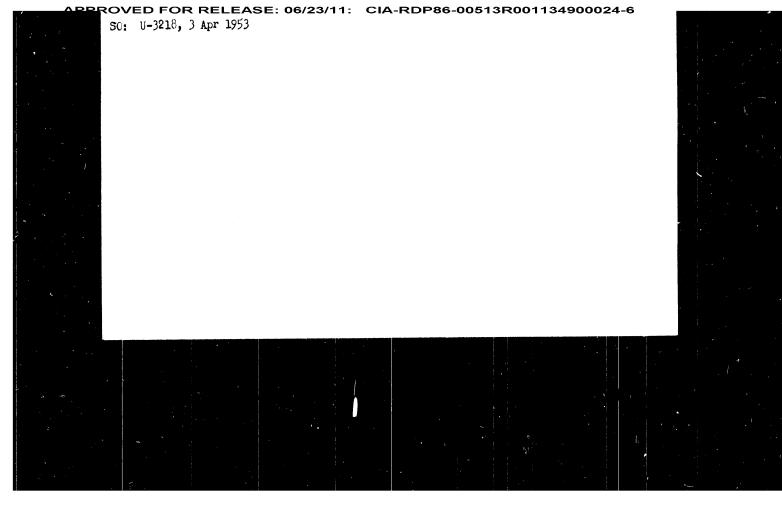


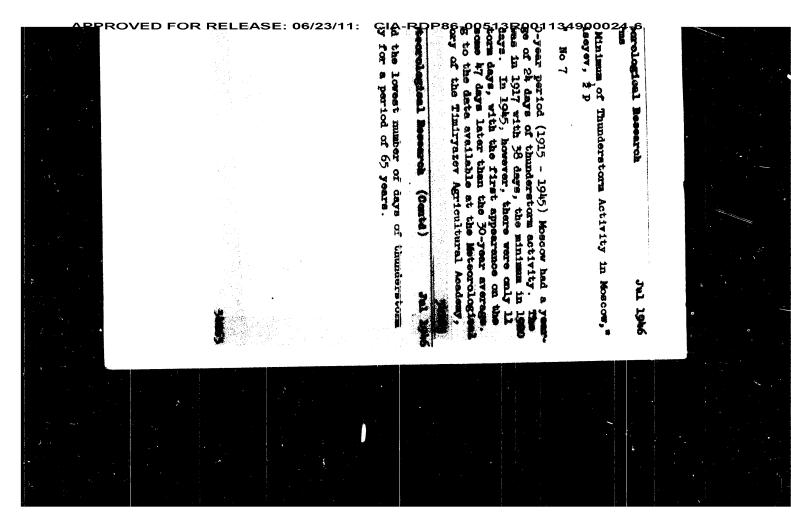


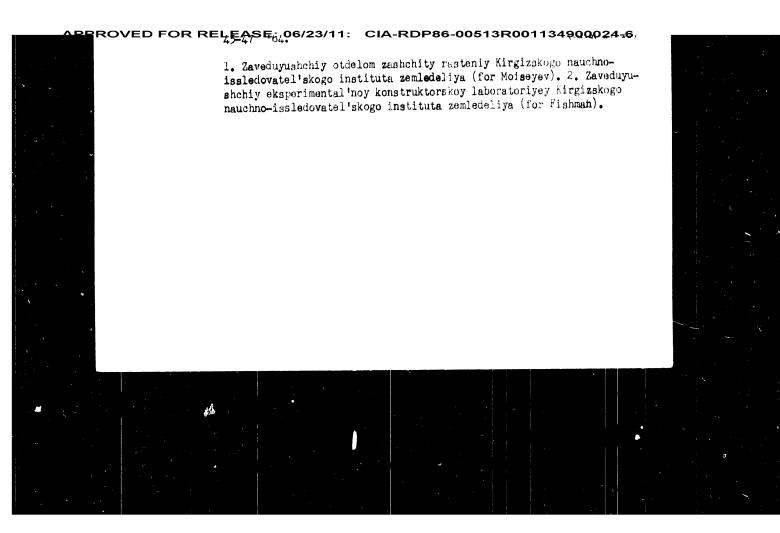












MOISEEN, A. N.

"Insects Which Danage Seeds and Fruit of the Woody Chrid Species of Activotians Oblact, Their Blobory and Control."

Cand Biol Sci. Novocherkands Coll Engineering Inst, Novocherkands, Coll Engineering Inst, Novocherkands, 1954. (Whitel, No. 6, Mar 55)

So: Sum No. 70, 29 Sep 55 - Survey of Schattific and Technical Assertations Defended at USSE, Higher Educational Institutions (15)

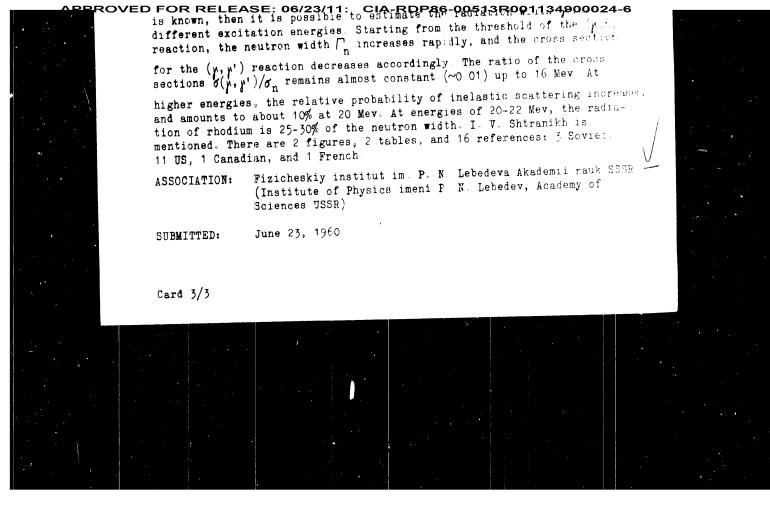
IVANOV, Yakov Andreyevich, kand. sel'khos. neuk, neuchnyy sotr.;
RYZERY, Ivan Petrovich, kand. biolog. neuk, neuchnyy sotr.;
ZAWOGAONYAYA, Yelena Tikhonovna, neuchnyy sotr.; TAPLOVA,
Yekaterina Alekseyevna, neuchnyy sotr.; MOISYEVI, Ilaksandr
Rikiforarich, neuchnyy sotr.; AEDUMANATOW, S., red.;
NOSOVETS, F.G., red.; HEYSHENOV, A., tekhm. red.

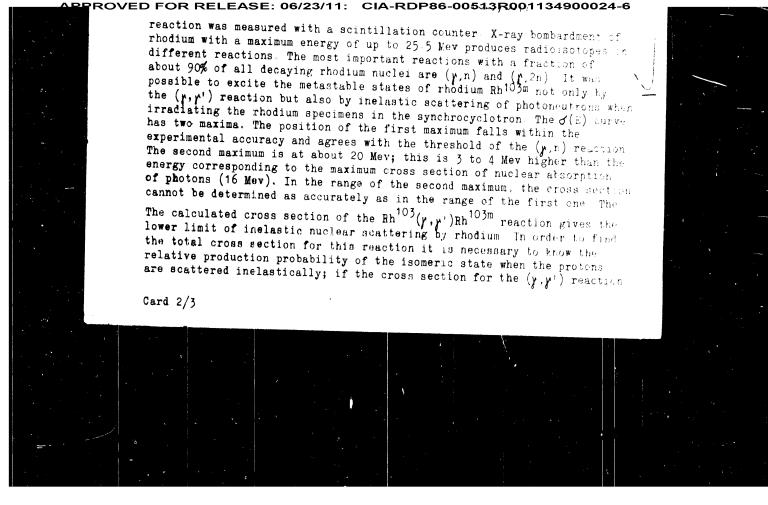
[Field testing of grain, oilseed, and forage crops in the
Kirghis S.S.R.) Aprobatsida zernovykh, mealichnych i kormovykh kul'tur v Kirgizskoi SSR. Frunze, Kirgizskoe izd-vo,
1959. 174 p.

[KIKA 15:3)

1. Kirgizskiy nauchno-issledovatel'skiy institut zemledeliya
(for Ivanov, Ryzhey, Zavgorodnyaya, Teplova, Moineyev).

(Kirghizistan—Orain breeding)
(Kirghizistan—Orain breeding)
(Kirghizistan—Forage plants)





AUTHORS:

Bogdankevich, O. V., Lazareva, L. Ye., Moiseyev, A. W.

TITLE:

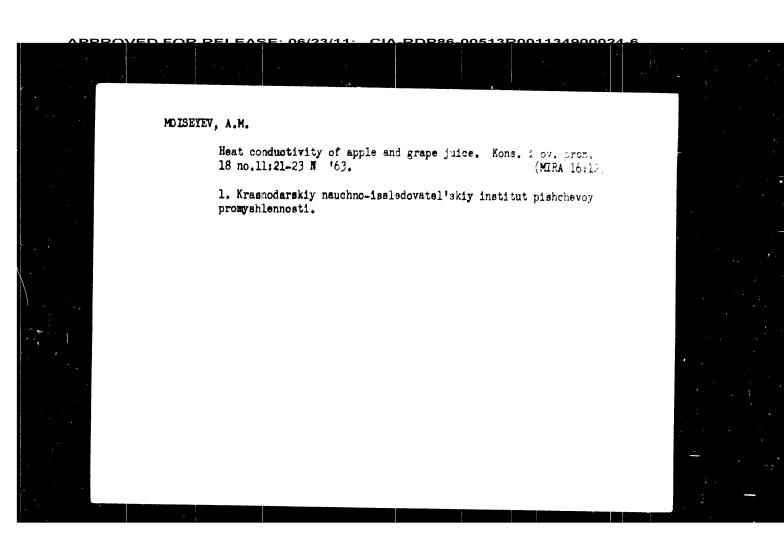
Inelastic Scattering of Photors by Rh 103 Nuclei

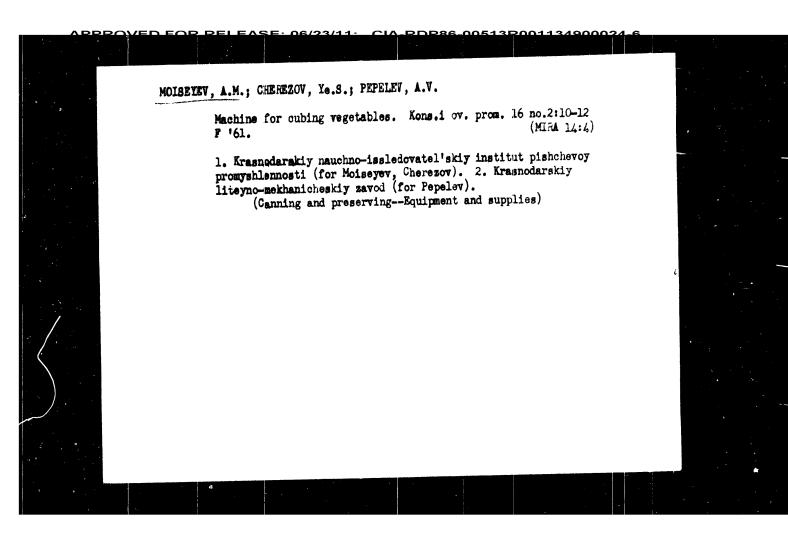
PERIODICAL:

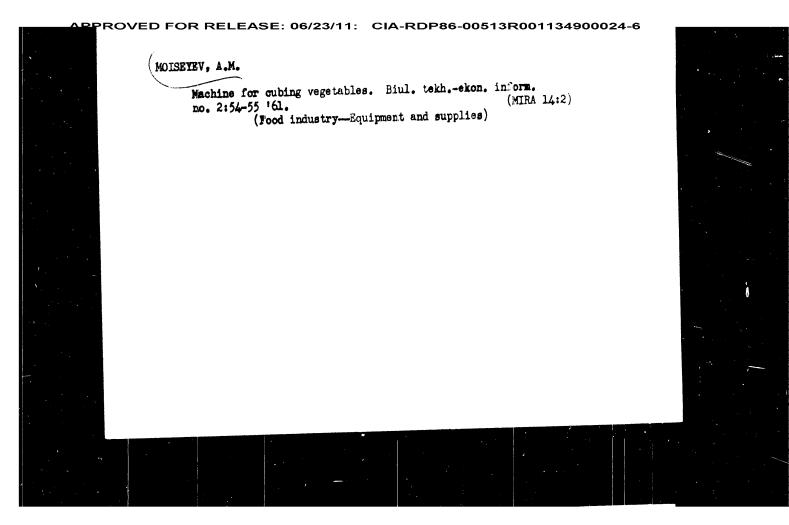
Zhurnal eksperimental noy i teoreticheskoy fiziki, 1962.

Vol. 39, No. 5(11), pp. 1224-1228

TEXT: The authors measured the yield of the reaction Rh 103 (y.y.) Rh 103m on the synchrotron of FIAN (Physics Institute of the Academy of Sciences in intervals of about 1 Mev. The number of isomeric nuclei of hh 103m induced activity. These measurements were made with specimens of metallic of Rh 103m nuclei is characterized by the following quantities: transition energy, 40 key; conversion coefficient a k from the K-shell, 40; ratio of the conversion coefficients on the K-shell and L-shell, 0.09 ± 0.01, and on the L-shell and M-shell, 7 ± 1. The yield of the Rh 103 (y.y.) Rh 103m



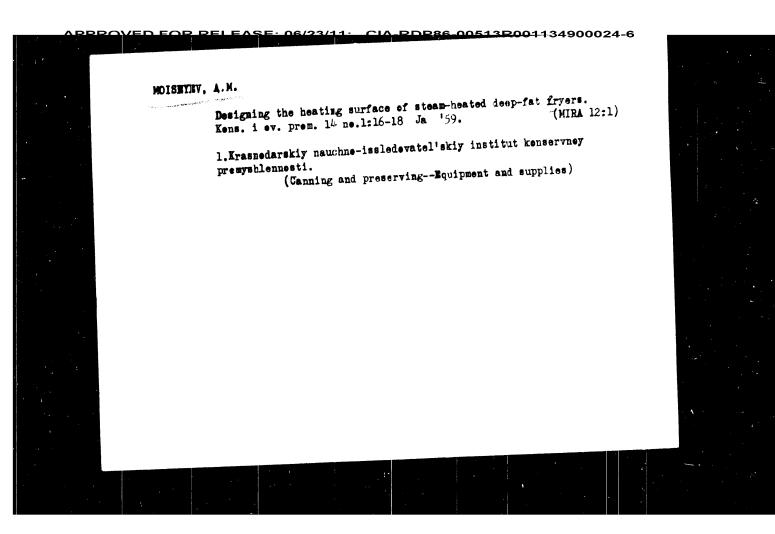


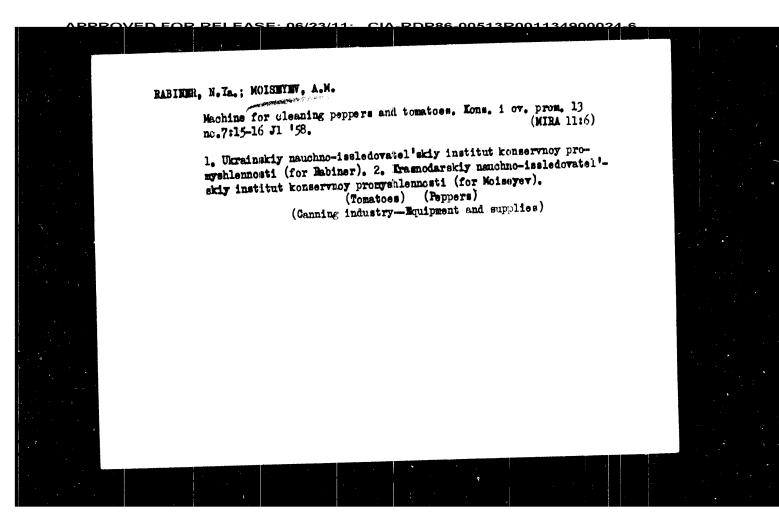


Moiserer, A.N.

Machines for peeling onions. Kons. i ov. prom. 14 no.8:10-12
Ag '59.

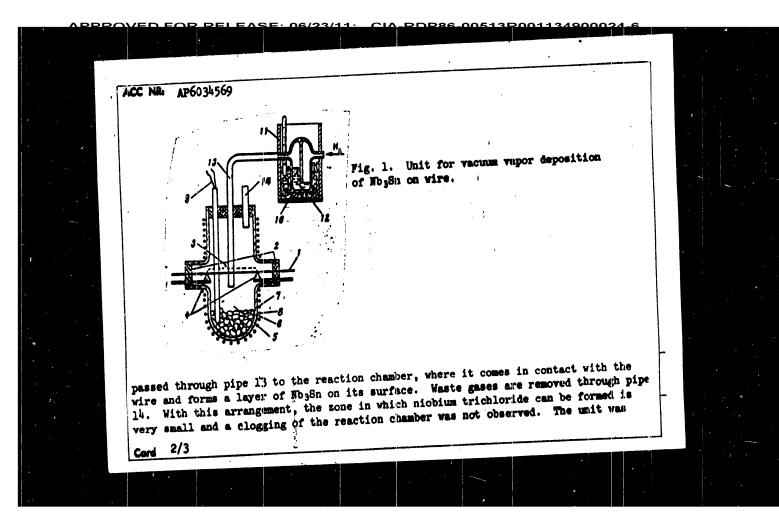
1. Krasnodarskiy nauchno-issledovatel'skiy institut pishchevoy
promyshlennosti.
(Onions)





tested on platinum and michrome wire 0.3 mm in diameter. The temperature of deposition was varied between 800—1200c. A single-phase layer (1—b p thick) of Mogan with B-W structure was obtained. It was established that the thickness of the layer increases with a temperature rise. Orig. art. has: 1 figure.

BUB CODE: 11, 13 SUBM DATE: 11May66/ ORIG REF: 003/ OTH REF: 013/
ATD PRESS: 5103



SOURCE CODE: UR/0020/66/170/006/1303/1305 (N) ACC NR. AP6034569 AUTHOR: Arkharov, V. I. (Academician AN Ukr&SR); Borisov, B. S.; Moiseyev, A. I.; Ugol'nikova, T. A. ORG: Institute of Physics of Metals, Academy of Sciences SSSR (Institut fiziki metallov Akademii nauk SSSR) TITLE: Vacuum vapor deposition of an No3Sn layer on a wire AN'SSSR. Dokelydy, v. 170, no. 6, 1966, 1303-1305 SOURCE: TOPIC TAGS: nightim time compound, superposition compound, sighting deposition, vacuum vapor deposition unti-ABSTRACT: To reduce the clogging of the reaction chamber by nonvolatile niobium trichloride, a new method and equipment (see Fig. 1) for continuous deposition of a superconducting layer of Mb₃Sn on a moving wire has been developed. Wire 1 is continuously fed through seals 2 into a reaction chamber at a fixed speed. Portion 3 of the wire is under treatment and is heated to about 1000C by electric current fed through sliding contacts 4. The bottom part 6 of reaction chamber 5 contains solid niobium pentachloride 7. The chamber is maintained at a temperature of 120-180° by electric furnace 8 controlled by thermocouple 9. Wepors of niobium pentachloride proceed directly to the wire. Hydrogen passing through reservoir 10, located in thermostat'll and kept at 0°C, is saturated with vapors of tin tetrachloride 12 and then UDC: 669.65.293:621.793 **Cord** 1/3

DORG: None

TITLE: Method of deposition of intermetallic nicbium-tin compound Nb₃Sn coating.

Class 48, No. 185661. [announced by the Institute of Physics of Metals, AN SSSR (Institut fiziki metallov AN SSSR)]

SOURCE: Izobreteniya, promyshlennyye obraztsy, tovarnyye znaki, no. 17, 1966, 128

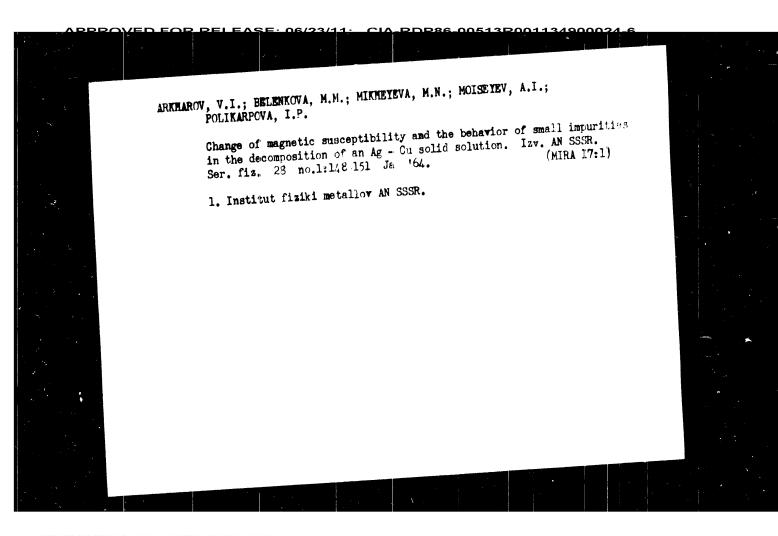
TOPIC TACS: niobium tin intermetallic compound, niobium tin compound coating, niobium tin compound deposition, NETAL DEPOSITION, METAL COATING, NIOBIUM COMPOUND, TON COMPOUND

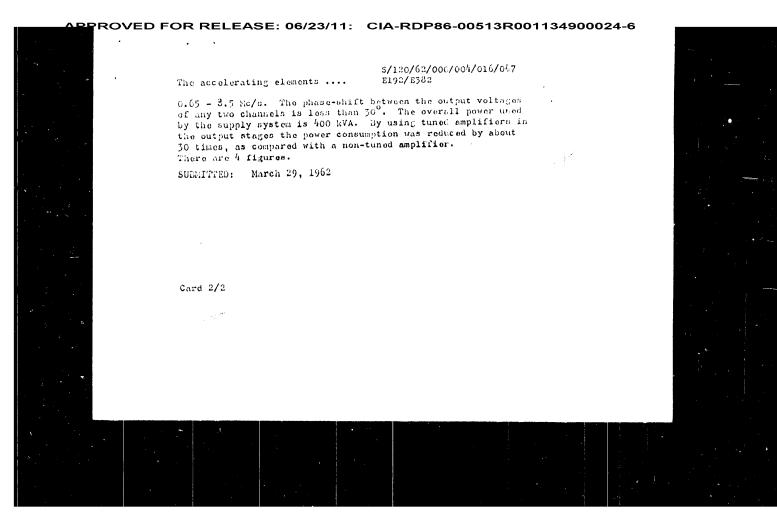
ABSTRACT: This Althor Certificate introduces a method of deposition of niobium tin compound coatings. To increase the purity and uniformity of the coating, niobium pentachloride is placed in the reaction chamber and heated to 120—160C, and the preheated port of the substrate.

SUB CODE: 11, 13/ SUBM DATE: 11Apr64/

Card 1/1 afs

UDC: 669.65' '293:621.793

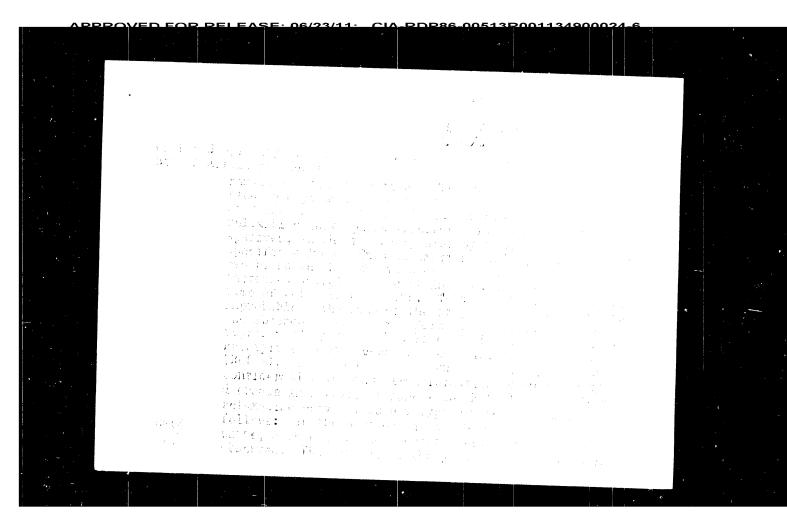


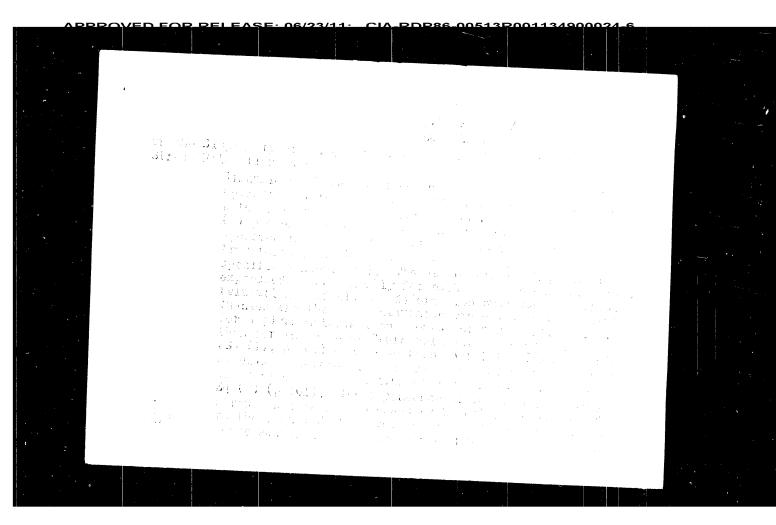


MOISEYEV, A. I. 1000 5/130/62/000/004/016/047 E192/E382 24.6036 Lobedov-Brasin, Yu.M., Gutner, B.M., Pisarevskiy, V.Ye., Teskin, A.S., Barabash, L.Z., Kurybaev, V.S. and holseyev, A.I. AUTHORE: The accelerating elements of the proton synchrotron and the system of their high-frequency feed TITLE: PORT ODICAL: Pribory i tekimika eksperimenta, no. 4, 1962, 94 - 97 The description, principal characteristics and the TEAT: results of the control of the h.f. accelerating system of the 7 GeV proton cyclotron are reported. The accelerating elements are in the form of drift tubes situated in 11 compensating . magnets. Each of the 11 electrodes is fed from a separate system of high-frequency amplifiers consisting of a 7-stage wideband amplifier and an automatically-tuned resonance output amplifier. The inductances of the resonant circuit in the output stages are in the form of coils fitted with ferrite cores. The amplitude of the high-frequency field of each accelerating electrode is 2.5 kV \pm 10% over the frequency range of Card 1/2

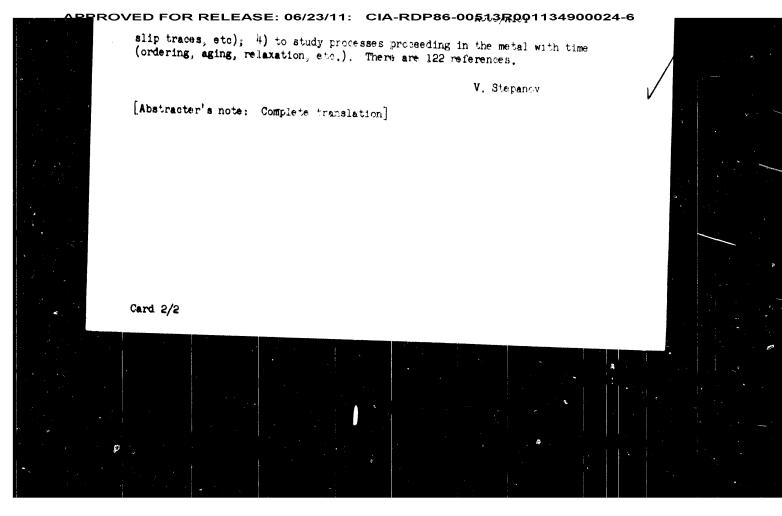
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Strong Simplicity is a Popular L Explanation of the person of the Simplicity is a Popular L Explanation of the person of the control deformation was considered as the factor as a considered by a description of the factor of the factor of the popular of the second of the strong of the person of the person of the factor of the second of t



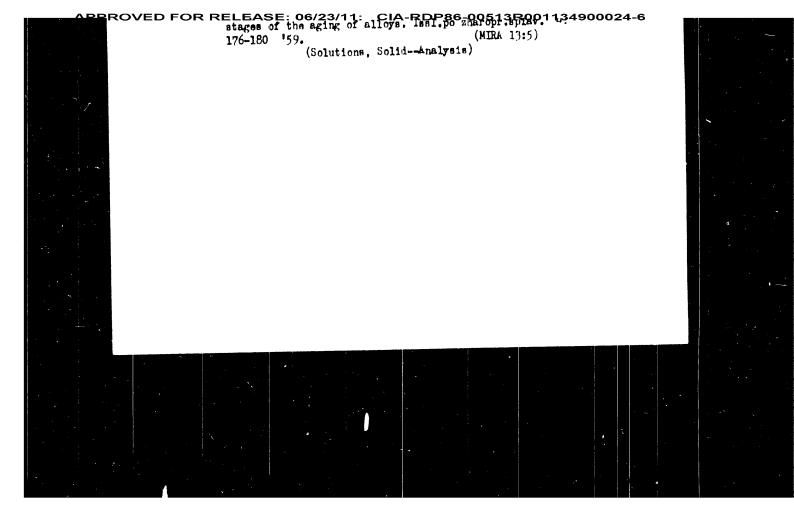


18 820. tilliffe. 8/336/60/009/03/03/03/03 BO(1/B1)1 AUTHORS: Kolesnikov, G.H., and Holsever, A.T. TITLE: On the Simplicity of a Physical Explanation of the Behaviour of Strees Relexating Ourves PERIODICAL: Finish manailes : manailes describe, 1960, v.a. o, so : ABSTRAST: In order to facilitate the argument the authors assume that the stress relaxation process occurs in a material which has been subjected to analytical tension. relaxation curve obtained experimentally under conditions of the so-called "pure relaxation" (Rof 1) represents the dependence of the average stress, acting on the specimen, on time, with the total deformation of the specimen being sept constant. As a result of imparting a deruain total deformation (i.e. new external conditions) to the specimen, various relaxation processes, different in their physical nature, arise in these houses. it which bring the specimen to a new equilibrium condition colles; causing to the coanged external conditions (Ref 2). These processes can be devided into Capyl. 1/5 two types: (1) processes causing an increase in



8/137/61/000/012/127/1-9 A006/A101 AUTHORS: Grin'; A.V., Moiseyev, A.I., Shmatov, V.T. TITLE: Internal friction and small amounts of admixtures in metals Referativnyy zhurnal, Metailurgiya, no. 12, 1961, 38, abstract PERIODICAL: 12Zh283 ("Tr. in-ta fiz. metallev, AN SSSR", 1960, no. 23, 163-173) This is a review. The authors analyze data-on the connection of internal friction with the presence of small amounts of admixtures. An analysis is made of internal friction in interstitional solid solutions; internal friction connected with relaxation along the grain boundaries, and internal friction caused by the presence of dislocations. The conclusion is drawn that with the aid of methods for the investigation of internal friction, it is possible; 1) to estimate the amount of small admixtures in pure metals, and in some cases, their concentration; 2) to determine diffusional parameters of atoms of admixtures at low temperatures; 3) to study internal adsorption of atoms of admixtures on structural heterogeneities (grain boundaries, domains, dislocations, Card 1/2

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SOV/126-6-5-40/43
An Internal Friction Peak Observed in the Testing of Deformed Aluminium

maximum B in curve 1 is due to relaxation of stresses at the points of localisation of plastic deformation. Tests of deformed aluminium (Ref 11) with temperature increasing show that the peak due to relaxation of stresses at the grain boundaries increases in magnitude but the second peak near 500°C does not appear at all. This is due to recrystallisation of the samples which occurs in the process of measurement itself, as the temperature is increased first to 300°C There are 1 figure and 12 references, 6 of which are Soviet and 6 English.

ASSOCIATION: Institut fiziki metallov AN SSSR (Institute of Metal Physics, Ac. Sc. USSR)

SUBMITTED: May 16, 1958

Card 3/3

An Internal Friction Peak Observed in the Testing of Deformed Aluminium

of the difference in the scales of the defects at the points of localisation of deformation and in the intercrystalline regions, the internal friction peaks, due to these two types of defects, should appear at different temperatures. To verify these ideas, the authors tested 99.0% alreading, compression-deformed by 80% by rolling and drawing down to 1 mm dia. Without intermediate annealing. Semplics were placed in a furnace previously heated to FAOC 3 and the internal friction was measured by season of a torsion pendulum (Ref 1). The internal friction was measured as the temperature in the furnace was lowered (Fig 1, curve 1). Curve 1 shows, in addition to the usual maximum A in the region of 500 C, an additional maximum B in the region of 500 C. Curve 2 of Fig.1 was obtained on aluminium samples which were deformed by 80% and then annealed for four hours at 600 C. Curve 2 has only one maximum A, which is due to relaxation of stresses in the grain boundaries. The suthors suggest that the

Card2/3

SOV/126-6-5-40/43 AUTHORS: Ibragimova, D.M., and Moiseyev, A. I. TITLE: An Internal Friction Peak Observed in the Testing of Deformed Aluminium (Pik vnutrennego treniya, nablyudayemyy pri ispytanii deformirovannogo alyuminiya) PERIODICAL: Fizika Metallov i Metallovedeniye, 1958, Vol 6, Nr 5, pp 952-953 (USSH) ABSTRACT: A peak in the region of 300°C (at 0.8 c/s frequency) was observed in the curve representing the temperature dependence of the internal friction in polycrystalline aluminium of 99.991% purity. It was first reported by Ke (Ref 1) and later confirmed by Ke (Refs 2-6) and by other authors (Refs 7-12). Ke ascribed the observed peak to relaxation of stresses at grain boundaries. He assumed that the grain foundaries (intercrystallite regions) have viscous properties, i.e. the crystalline structure in the intercrystallite regions is defective. The present authors point out that deformed metals have a large number of points of localisation of plastic deformation. Viscous properties of the points of localisation of deformation should also affect the Cardl/3 temperature dependence of the internal friction. Because The Effect of Small Additions of Antimony and Beryllium on Ageing of the Copper-Silver Alloys. (Part 1V. On the Problem of Causes of the Effects of Small Alloying Additions on the Kinetics of Ageing of Alloys)

of the lattice distortions in the zones connecting the nuclei with the solid solution matrix are correspondingly altered. There are 9 graphs, 1 table and 21 references of which 20 are Soviet and 1 English.

ASSOCIATION: Institut Fiziki Metallov Ural'skogo Filiala AN SSSR (Institute of Metal Physics, Ural Branch of the AS USSR SUBMITTED: 18th December 1956.

807/126-6-4-9/3-

The Effect of Small Additions of Antimony and Beryllium or keeping of the Copper-Silver Alloys. (Part 1V. On the Problem of Causes of the Effects of Small Alloying Additions on the Kinatics of Againg of Alloys)

vicinity of the solute atoms are non-additive because - owing to the fact that Be atoms are smaller and the atoms larger than the solvent atoms - the lattice distortions caused by the atoms of either element present separately are more severe than those caused by the complexes formed when the two alloying additions are present simultaneously. In the later stages of the ageing process when growth of the decomposition centres affected by the adsorption of the alloying elements in the surrounding zones is the predominating factor, the non-additive character of the effects of 56 and 28 is evidently due to the fact that at first beryllium is preferentially adsorbed, while adsorption of antimony takes place mainly in the later stages. This time-19g in the adsorption activities of the two elements is probably associated with the fact that with the growth Card 10/11 of the decomposition nuclei the character and magnitude

20V/1256 --- 9/3

The Effect of Small Additions of Antimony and Beryllium on Ageing of the Copper-Silver Alloys. (Part 1V. On the Problem of Causes of the Effects of Small Alloying Additions on the Kinetics of Ageing of Alloys)

overall concentration of the additions present simultaneously in the alloy: At a given Sb concentration, beryllium - when present in small quantities - is absorbed mainly in the form of compasses with the result that the concentration of Sb in the adsorption zone is increased and its accelerating effect on the decomposition of the solid solution is multiplied. On the other hand, when the Be content is high, it is adsorbed in the form of single atoms which increases its concentration in the adsorption zones with the result that the rate of decomposition is slowed down. The effects of Be and Sb on the course of the againg process are also non-additive. In the initial stages of the process when formation of nuclei of decomposition is the predominant factor affecting the kinetics of decomposition, the effects of the alloying additions on nucleation due to local lattice distortions in the

Card 9/11

SOV/126-E-4-9/34

The Effect of Small Additions of Antimony and Beryllium or Ageing of the Copper-Silver Alloys. (Part IV. On the Problem of Causes of the Effects of Small Alloying Additions on the Kinetics of Ageing of Alloys)

including those corresponding to the maximum values of vcm. These effects can be explained on the basis of a hypothesis of internal adsorption of the So and Be atoms in structurally distorted zones linking the nuclei of decomposition with the solid solution matrix: it being postulated that the alloying elements can be adsorbed not only as separate aroms but also in the form of complexes containing atoms of both additions. As a result of the adsorption of complexes the free energy of the distorted zones is decreased in regions where owing to the specific character of the distortion - it would not be decreased by adsorption of single atoms. The extent to which adsorption of complexes affects the kinetics of decomposition of the solid solution varies with time since, as a result of adsorption, the total concentration of both alloying elements in the adsorption zone is altered to a degree depending on the

Card 8/11

SOV/126-5-4-9/34

The Effect of Small Additions of Antimony and Beryllium on Ageing of the Copper-Silver Alloys. (Part 1V. On the Problem of Causes of the Effects of Small Alloying Additions on the Kinetics of Ageing of Alloys)

stage, its effect being most pronounced at t = 50 min. The following interpretation of the obtained results is offered by the present authors: The average rate v_{cm}, of the isothermal decomposition of a supersaturated solid solution of silver in copper is markedly affected by small simultaneously present additions of Sb and Be, even when these elements are present in concentrations considerably lower than their respective solid solubility limits. When added separately, antimony accelerates and beryllium slows down the process of decomposition. However, these effects are not additive when Sb and Be are present simultaneously: At a given Sb concentration vom increases at first with the increasing Be content reaches a maximum and then slowly decreases (Fig.1-3). The higher the content of antimony the higher are the Card 7/11 values of vom for any given beryllium concentrations

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SOV/126-6-4-9/34

The Effect of Small Additions of Antimony and Beryllium on Ageing of the Copper-Silver Alloys. (Part 1V. On the Problem of Gauses of the Effects of small Alloying Additions on the Kinetics of Ageing of Alloys)

and the greater is the magnitude of the effect.) The effect of 0.2% Sb on againg of Cu-Ag alloys containing 0.02 and 0.1% Be (Fig.5) is similar to its effect on the binary Cu-Ag alloy. The same applies to the effect of simultaneous additions of Sb and Be, except that in this case the maximum value of Ax decreases with increasing Be content (Fig.6). The effect of Be on kinetics of ageing of the Cu-Ag alloy containing 0.2% Sb is much more complex. At small concentrations (0.02%) beryllium accelerates ageing of the Cu-Ag-Sb alloy (graph 1) in all stages of the process, Ax reaching its maximum after 1 hr. 0.1% Be slows the process down in its initial stage and accelerates it slightly in the final stage. When present in larger quantities (0.2 - 0.3%) it slows down the ageing process of the Cu-Ag-Sb alloy at every

Card 6/11

SOV/126-6-4-9/34

RDP86-00513R001134900024-6

The Effect of Small Additions of Antimony and Reryllium on Ageing of the Copper-Silver Alloys (Part 1V. On the Problem of Causes of the Effects of Small Alloying Additions on the Kinetics of Ageing of Alloys)

property to the length of time required to effect this variation) was calculated for various investigated alloys and the results were reproduced graphically. Figl shows how vcm (assessed on the basis of hardness measurements) of alloys with a constant Sb content aged at 370°C varied with increasing Be content. The variation of vcm (calculated from the data on magnetic susceptibility) of alloys containing 0.2% Sb and aged at 370, 400 and 420° with increasing Be content is shown in Fig.2, while Fig.3 shows the effect of Be on vcm (determined on the basis of electrical resistance measurements) of the 0.2% Sb alloy aged at 370°C. The effect of the Sb and Be additions on the course of the ageing process in its various scages was determined on the basis of the measurements of magnetic susceptibility, since this property could be measured

Card 4/11

80V/126-6-4-9/3-

The Effect of Small Additions of Antimony and Beryllium on ageing of the Copper-Silver Alloys (Part 1V. On the Problem of Causes of the Effects of Small Alloying Additions on the Kinetics of Ageing of Alloys)

15-30 minute intervals on specimens solution treated at 780 - 790°C and aged at 370°C. Magnetic susceptibility was measured with the aid of a magnetic balance at room temperature and at 370, 400 and 420°C. The measurements were taken at 10-15 minute intervals and in every case the value of relative magnetic susceptibility was determined, i.e. the force acting on the investigated specimen was compared with the force acting on a standard nickel sulphate specimen placed in an identical magnetic field. Electrical resistance was measured by the comparison of potential drop method, using a potentiometer and a sensitive galvanometer. In this case, both the solution treatment and ageing (at 370°C) were carried out in vacuum and the measurements were taken at 15 minute intervals. From the experimental data the average rate of ageing (v_{cm} = the ratio of the maximum increment of the studied

Card 3/11

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SOV/126-6-4-9/34

The Effect of Small Additions of Antimony and Beryllium on Ageing of the Copper-Silver Alloys (Part IV. On the Problem of Causes of the Effects of Small Alloying Additions on the Kinetics of Ageing of Alloys)

additions. The object of the present investigation was to study the simultaneous effect of two horophilic additions. The experimental alloys whose detailed chemical analysis is given in a table on p 633, contained 6% Ag with 0.2 - 0.5% Sb and 0.02 - 0.3% Fe added either separately or jointly. The alloys were melted in a H.F. induction furnace, in a graphite crucible with corax used as the covering flux. The cast ingots were heated under charcoal to 80000 held at the temperature for 2 hrs and cooled in the furnace. They were then relled to strip 5 mm thick which, after a homogenising treatment consisting of 50 hours at 800°C was used for the preparation of the experimental test pieces. The process of ageing was studied by measuring the variation of hardness, magnetic susceptibility and electrical resistance. The Card 2/11 measurements of Rockwell hardness were taken at

MOISEYEV, A. I.

- with V. S. AVERKIYEV, G. N. KOLASNIKOV, and M. V. YAKOTOVICE "Device for Testing of Stress Relaxation in Tencion" p. 71
- with G. N. KOLESNIKOV and M. V. YAKUTOVICH
 "Effect of Small Additions of Alloying Elements on Stream Remarkle For Iron-Chrome-Nickel Alloys" p. 95
- with G. N. KOLESNIKOV
 "Slip Plasticity and Diffusion Plasticity in the Streen-Relaxation Process"
 p. 101
- with V. I. ARKHAROV, S. I. IVANOVSKAYA, G. N. KOLESNIKOVA
 "Stress Relaxation and Irregularity of Liffusional Mobility in
 Polycrystalline Austenitic Iron-Chrome-Nickel Alloys" p. 1823
- with G. N. KOLESNIKOV "The Effect of Phase Transformations on Stress Relaxation" p. 107

Problems in the Theory of Heat Resistance of Metal Alloys, M. -w., 12)-77
AN SSSR, 1958, 160 pp. (Trudy, virt. Fiz. Metal., Ural fillal, AL SSSR)
76, 17

The articles in this book constitute reports on extensive studies, nondrotes between 1949 and 1954 by the Inst. Physical Metallargy Urals Branch, AS USSE, and devoted to the development of a general theory of heat resistance.

SOV/137 58 8 17694

An Investigation of the Effect of Small Quantities (cont.)

solution. Owing to adsorptional concentration changes in these transitional zones, the rate of Ag diffusion, which is instrumental in the expansion of primary zones of increased density and in further growth on newly formed crystal nuclei of the precipitating phase, is greatly changed. Bibliography:

18 references.

V. N.

1. Antinony-ber/llium-copper-cilium elloyes—
Analysis 2. Antimony-beryllium-copper-cilium alloys—lging

Card 3/3

APPROVED FOR RELEASE: 06/23/11: CIA-RDP86-00513R001134900024-6

SOV /117 58 8 17694

An Investigation of the Effect of Small Quantities won't

charcoal packing for a period of 50 hrs at a temperature of 800° and were allowed to cool in air. After a two hour exposure to 780 7900 the specmens were quenched in ice water. The A was carried out at 300° first stage of the process (up to maximum hardness of the afloy) requiring from 0.5 to 2 hrs; the total duration of the process amounted in 3.5 krs. Every 15 minutes the specimens were taken out of the turnace and were cooled in water, after which their hardness was measured. The average rate of aging was determined from the curves showing the hardness as a function of the A time. It was established that the accelerating and retard ing time rate-affecting additives, present concurrently in a solid section undergoing decomposition do not have an additive effect upon the process of A. A maximum rate of A was obserted translatory with a minimum an outof Be additive (in the presence of Sb); the absolute magnitude of they rate is greater than the rate of A in an alloy with the same concentration of Sh but containing no Be. The absolute magnitude of hard ess of alloys containing both Sb and Be is greater than that of alloys containing it add these or only one of the additives. The strong influence of small amounts of Sb and Be on the rate of A is explained by internal adsorpt on of Sh and Be in distorted zones of the junctions of the primary regions with increased density of Ag concentration, or in zones of new phase nuclei with their surrounding solid Card 2/3

SOV/137-58-8-17694

Translation from: Reterationyy zhurnal, Metallurgija (1948 Nr. 8 5211/USSR)

AUTHORS: Arkharov, V. I., Moiseyev, A. L., Polikarpora, I.P.

TITLE: An Investigation of the Effect of Small Quantities of Additives on

the Kinetics of Aging of Alloys (Issledo: aniye ·livaniya malykh

primesey na kinetiku stareniya splavo)

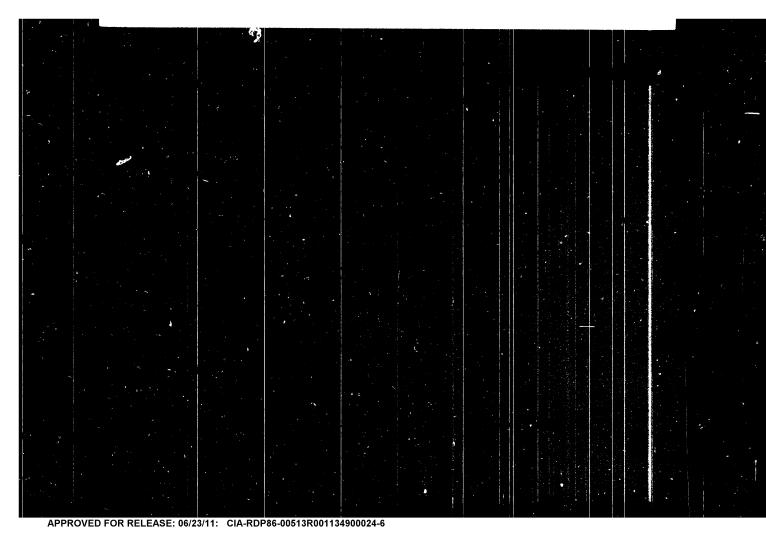
PERIODICAL: V sb.: Issled po zharoprochu splavam Vol 2 Moscow, AN

SSSR 1957 pp 92 97

ABSTRACT: Hardness measurements were employed in an investigation dealing with the combined accelerating and retarding effect of time-rate affecting additives on the early stages of the aging (A) process of Cu alloys containing up to 6% of Ag. Sh (up to 0.5%) and Be (up to 0.3%) served respectively as the accelerating and retarding additives. The alloys were prepared from electrolytic Cu (99.9% pure), Ag and Sh (both 99.9% pure), and Be (97.7% pure). Following a two hour anneal at a femperature of 800°C and a period or cooling in the turnace, the ingots were forged at room temperature, in order to reduce

their thickness from 8 mm to 6mm and relled until their final thickness amounted to 5 mm; they were then homogenized in a

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MOISEYEV, Aleksey Grigor yevich; PETROV, Viktor Mikbaylovich; VOLKOV, I.V., retsenzent; VERBITSKAYA, Ye.M., red.; SHUB, L.S., spets.red.; SHVETSOV, S.V., tekhn.red. [Manual for engraving of textile patterns] Rukovodstvo po gravirovamiu tekstil'nogo risunka. Noskva, Izd-vo nauchno-tekhn.lit-ry RSFSR, 1961. 147 p. (MIRA 15:2) (Textile printing) (Engraving)

Control of the Contro

ACCESSION NR: AP4043823

dipped parts were exposed for 10 days to sea water or nitric acid (\sim 17% concentration). Prior phosphating did not improve the corrosion resistance of coatings. FG-9, FG-50 and MK-4 were best in tap water, FG-9 and FG-50 were adequate in 5% sodium chloride at 50C and 5% solutions of nitric or hydrochloric acids, but not in sulfuric acid. MK-4 protected only in dilute solutions of nitric acid. Adhesion to an untrested scale surface was poor for all coatings. Two coats of FG-9 or FG-50 provided adequate protection in sea water. FG-50 was somewhat better in nitric acid and is recommended. The stability of its film increases when curing temperature is increased to 200C. Orig. art. has: 5 tables.

ASSOCIATION: None

SUBMITTED: 00

ENCL: 00

SUB CODE: MT

NO REF SOV: 000

OTHER: 000

ACCESSION NR: AP4043823

8/0303/64/000/004/0054/0056

AUTHOR: Moiseyev, A. F., Olenin, S. S., Neshumova, A. I.

TITLE: Use of polyorganosiloxane coatings to protect parts of DM-6 differential manometers against corrosion in aggressive media

SOURCE: Lakokrasochny vo materialy i ikh primeneniye, no. 4, 1964, 64-56

TOPIC TAGS: anticorrosion coating, polyorganosiloxane coating, differential manometer, coating FG-9, coating 192T, coating FG-50, coating MK-4, coating FG-35, chromic oxide coating, aluminum powder coating, titanium dioxide coating, metallic coating, phosphate precoating, polyorganosiloxane, corrosion, sea water

ABSTRACT: Modified polyorganosiloxane coatings 192T, FG-9 (+ 12% chromic oxide or 6% Al powder by weight), FG-35 (+ 5% Al powder), FG-50 (+ 5% Al powder) and MK-4 (+ 12% titanium dioxide) were used to coat the housings, seats and membrane chambers of DM-6 differential manometers in a study of optimal protection against corrosion in various aggressive media. Housings and seats were prepared by sandblasting, membrane blocks were either untreated or etched in nitric-hydrochloric acid mixture. Sprayed parts (1 or 2 coats) were exposed for one month to tap water at room temperature, 5% sodium chloride at room temperature or 50C, or 5% solutions of sulfuric, hydrochloric or nitric acids;

NADZHAFOV, Yu.B.; LOSEV, V.B., DHAUTOV, Yu.Kh., MOINFEFV V.J. TUBYANSKAYA, V.F. Heats of combustion of some mitrogen-containing organ is some Thur. fiz. khim. 39 no.5:1220-12.3 My 188. The Park Fig.

L 1801:3-66 ACC MR: AP6004314 use was made of disethylchlorosilane, methylphenyldichlorosilane (MP), phenyltrichlorosilene, and methyltrichlorosilene (M). Comparative studies of polyorganocharylance licrame and polyorganosiloxane lacquer films showed that both the polyconensatice and polymerisation of cohydrolymetes of organophenylenesiloxanes occur much Easter (5 to 15 times) than those of cohydrolymates of organosiloxanes. Because of replacement of a large number of siloxane linkages in the main chain of polyorganosilements by silphenylene linkages, the drying time of the films is substantially reduced and their thermal stability is improved, but their thermoelasticity selimen (i. a., mittleness increases). Introduction of small quantities of silinse linkages into the polyorgenosiloxenes, carried out by adding small amounts of p-bis (phenyldich leroeily 1) beasens and bis (methyldichloroeily 1) beasens to the initial sixture of chlorosilanes, was shown to shorten the drying time of the films, increase their mechanical strength and thermal stability, and leave the thermal stability and thermoslasticity unaffected. Orig. art. has: 2 figures, 4 tables. ORIG REF: 009/ OTH REP: 012 SIBM DATE: 00/

APPROVED FOR RELEASE: 06/23/11: CIA-PDP86-00513P001134900024-6

1 10012-66 EWT(x)/ENP(j) RM

SCURCE CODE: UR/0303/65/000/005/0014/0018

ATTHOR: Molecyer, A. F.

ORG: name

TITLE: Synthesis and some properties of lacquer films from polyorganosiloxame poly-

SOURCE: Lakokresochnyye materialy i ikh primeneniye, no. 5, 1965, 14-18

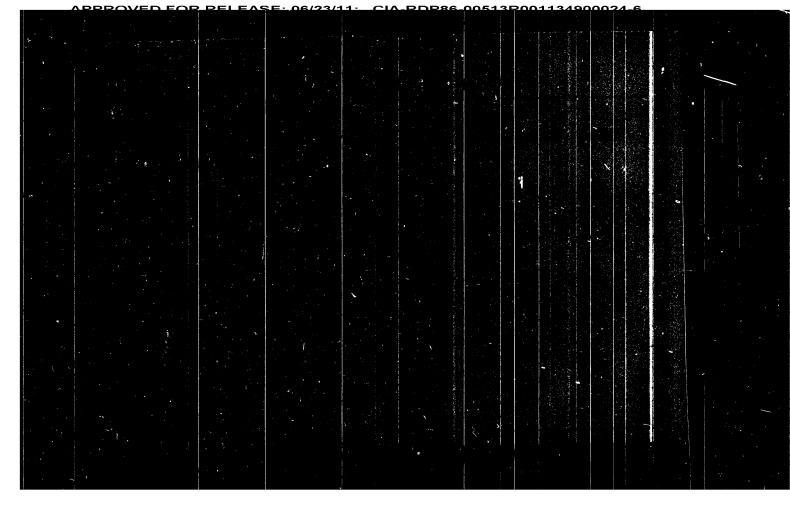
TOPIC TAGS: polymer structure, polysiloxane, organosilicon compound

ABSTRACT: A study was made of the influence of the nature of the organic solvent on the course of onlydrolysis of various bis[alkyl(aryl)chlorosilyl]benzenes and alkyl (aryl)chlorosilanes, and of the influence of conditions of cohydrolysis and relative propertions of these chlorosilanes (functionality of the mixture), on the polycondensing (polymerizing) capacity of the cohydrolysates and on the properties of the cohydrolysates obtained, i. e., polyorganophenylanesiloxanes. The phenylane-containing memorars used were bis(trichlorosilyl)bensene, bis(methyldichlorosilyl)bensene, and p-bis(phenyldichlorosilyl)bensene (BPSB). Among alkyl(aryl)chlorosilanes,

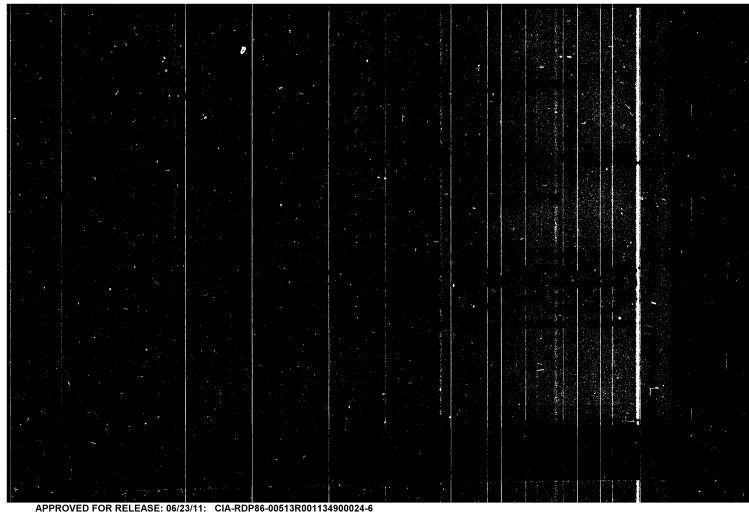
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26992

Production of linear polyurethanes...

S/191/61/000/009/002/007 B110/B218

affected by the diol ratio. There are 7 figures, 5 tables, and 6 references 2 Soviet and 3 non-Soviet. The two references to English-language publications read as follows: Ref. 1: 0. Bayer, Mod Plastics, 24, 149 (1947); Ref. 5: R. Hill, E. Walker, J. Polymer Sci., 3, 609 (1948).

Table 3. Physicomechanical properties of PU obtained from hexamethylene diisocyanate and glycol mixtures. Legend: (1) glycol, (2) glycol ratio, moles, (3) impact strength according to Dinstat, kg·cm/cm², (4) static bending according to Dinstat, kg·cm/cm², (5) Brinell hardness, kg/mm², (6) water absorption, %, (7) thermal stability according to Vicat, °C, (8) melting temperature, °C, (9) ethylene glycol, diethylene glycol, (1C) ethylene glycol, 1,4-butanediol, (11) diethylene glycol, 1,4-butanediol, (12) triethylene glycol, 1,4-butanediol, (13) 1,6-methyl hexanediol, 1,4-butanediol.

Card 4/5

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Production of linear polyurethanes...

S/191/61/000/009/002/007 B110/B218

films. Melting point, thermal stability, Brinell hardness first drop with increasing EG content of the mixture, start rising at 35-40% of EG and reach a maximum at 100% of EG. The water-absorbing capacity drops to 0.23% with 100% of EG. PU on the basis of an EG - 1,4-butanediol (BD) mixture are white, horny, but less solid, soluble only in cresol and a phenol-water mixture (90: 10). In melting, they yield strong threads and elastic films. Melting temperatures, thermal stabilities, and impact strengths are lower in glycol mixtures than in individual diols, the water-absorbing capacity is slightly higher. PU on the basis of DE - 1,4-BD mixtures are transparent, elastic, and soluble in cresol only. The elasticity of threads and films from the melts decreases with increasing 1,4-BD content while the strength increases. PU on the basis of triethylene glycol - 1,4-BD mixtures are more elastic, and soluble in $C_2H_5OH-CHCl_3$ and $C_2H_5OH-C_2H_4Cl_2$. In the same soluble vents, also PU based on methylhexanediol- 1,4-BD mixtures are soluble. These PU are white or transparent elastic products. From the melt, they produce very strong threads and solid films. The impact-strength curve

according to Dinstat of the DE - 1,4-BD mixture passes a minimum with increasing DE content, and reaches a maximum at about 75% of DE. The physicomechanical properties of PU are strongly and in a complicated manner

Card 3/5

26992

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Production of linear polyurethanes

threads in melting. They are soluble in trioresol or in a phonoi-water mixture (90: 10). PU based on diethylene glycol and 1,6-methylhexanedical are transparent colorless elastic substances; they are soluble in tricresol, in phenol-water mixtures (90: 10), and also in alcohol-chloreform mixtures (50: 50). From the solutions, they form elastic films of high mechanical strength. Transparent, brittle products are obtained on the basis of glycols with a secondary hydroxy group such as 1.2-propanecial and 1,3-butanediol. Pu hardness drops with rising molecular weight of diol The side groups and the oxygen atom in diol increase the impact strength PU on the basis of diethylene glycol (DE) have low strength and thermal stability, high water-absorbing capacity, and a large tangent of the angle of dielectric losses. The authors found: The presence of a methyl side group or of ether oxygen in the glycol component reduces the melting temperature, hardness, thermal stability, and other physicomechanical indices. Elasticity and solubility in organics increase. PU on the basis of an ethylene-glycol (EG) - DE mixture are white, elastic, soluble in crescl, $C_2^{\prime}H_5^{\prime}OH \sim CHCl_3^{\prime}$ mixtures (50 : 50), and $C_2^{\prime}H_5^{\prime}OH \sim C_2^{\prime}H_4^{\prime}Cl_2^{\prime}$ From melts, PU with high EG content produce strong threads and poorly elastic films, PU with high DE content, weak threads and solid elastic

APPROVED FOR RELEASE: 06/23/11: CIA-RDP86-00513R001134900024-6

3/191/61/000/009/002/00 26992 B110/B218 15.8140 Korshak, V. V., Strepikheyev, Yu. A., Moiseyev, A. F AUTHORS. Production of linear polyurethanes without solvents. Some TITLE: physicomechanical indices of polyurethanes based on hexamethylene diisocyanate and a number of glycols Plasticheskiye massy, no. 9, 1961, 16 - 20 PERIODICAL: TEXT: The authors studied the change in physicomechanical properties of polyurethanes (PU) obtained on the basis of hexamethylene discovanate and some glycols. The PU were produced by adding the stoichiometric amount of dissocyanate to the heated diol. The method was developed by the authors (Plast. Massy, No. 6, 1961, ibid., No. 7, 1961). The authors determined melting point, molecular weight (viscosimetrically), and other physico. mechanical indices. In PU on the basis of diols with different numbers of C atoms, the maximum melting point lies at 18300 for 1,4-butanediol PU. Melting points of PU on the basis of glycols with even numbers of C atoms are slightly higher than with odd numbers of C atoms. Pu on the basis of ethylene glycol, 1,4-butanediol, 1.6-hexanediol, and 1,10-decanediol are transparent or white, solid, horny substances which yield sold-drawing Card 1/5 APPROVED FOR RELEASE: 06/23/11: CIA-RDP86-00513R001134900024-6

24745

\$/191/61/000/007/003/010 B101/B215

Synthesis of linear polyurethanes...

ROVED FOR RELEASE: 06/23/11:

weight, the Staudinger constant was set equal to $11 \cdot 10^{-4}$. 1) In the presence of oxygen, the molecular weight was considerably reduced. In nitrogen atmosphere, the molecular weight was only slightly reduced (from 36,000 to 33,000) by an increase of the initial temperature of the reaction from 80 to 180°C . In the presence of air, the molecular weight was 25,000 at 80°C and 14,000 at 180°C. An increase of the final reaction temperature from 190 to 230°C reduced the molecular weight in nitrogen atmosphere from 36,000 to 23,000 and in air from 24,000 to 8000. Under optimum reaction conditions (initial temperature: 80 to 90°C; final temperature: 200-210°C) the duration of dropwise addition of diisocyanate in nitrogen atmosphere did not affect the molecular weight. By dropwise addition of 20 min, a maximum molecular weight of 25,000 was obtained in the presence of air. Slower addition reduced the molecular weight. After 5 hr the molecular weight was unchanged by heating the polymer in N2 up to 200°C. Heating up to 210°C led to a slight decrease. The presence of air reduces the molecular weight by 50 % within 2 hr at 190°C or within 1 hr at 200°C. 2) An excess of 1,4-butanediol caused the following changes in the molecular weight: 0.0 % excess: 35,400; 1.0 %: 28,050; 10 %: 5100;

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AUTHORS:

Korshak, V. V., Strepikheyev, Yu. A., Moiseyev, A. F.

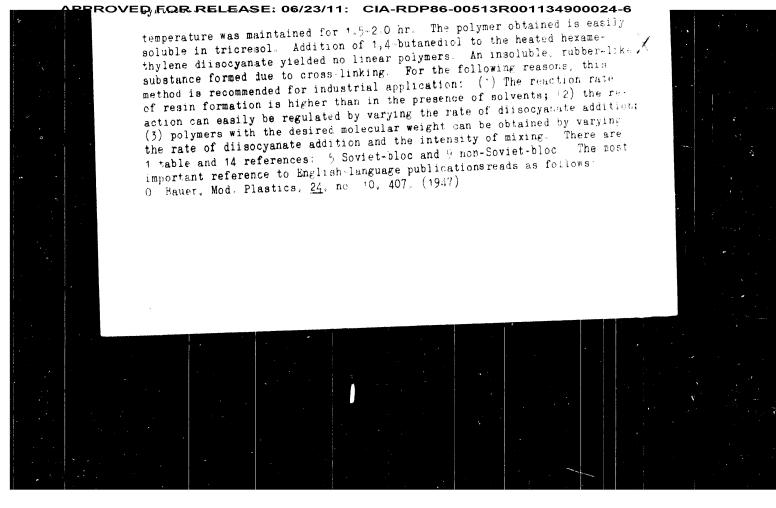
TITLE:

Synthesis of linear polyurethanes without solvents. Some rules governing the reaction of hexamethylene diisocyanate with 1,4-butanediol in the melt

PERIODICAL: Plasticheskiye massy, no. 7, 1961, 13-16

TEXT: This report is a continuation of a study on the synthesis of polyurethanes without using solvents (Plast. massy, no. 6, 1961). The effect of the following factors upon the molecular weight was studied in the present paper: 1) presence of atmospheric oxygen; 2) ratio of the components; 3) addition of monofunctional reagents; 4) duration of heating. The synthesis of polyurethane was conducted by dropwise addition of hexamethyl disocyanate to heated 1,4-butanediol. The temperature of the mass was not allowed to exceed a certain reaction temperature. The melt was kept at this temperature for a while, and, finally, the viscosity of a 0.5 % solution in tricresol was determined. In calculating the molecular

Card 1/3



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S/191/61/000/006/002/005 B101/B215

AUTHORS:

Korshak, V. V., Strepikheyev, Yu. A., Moiseyev, A. F.

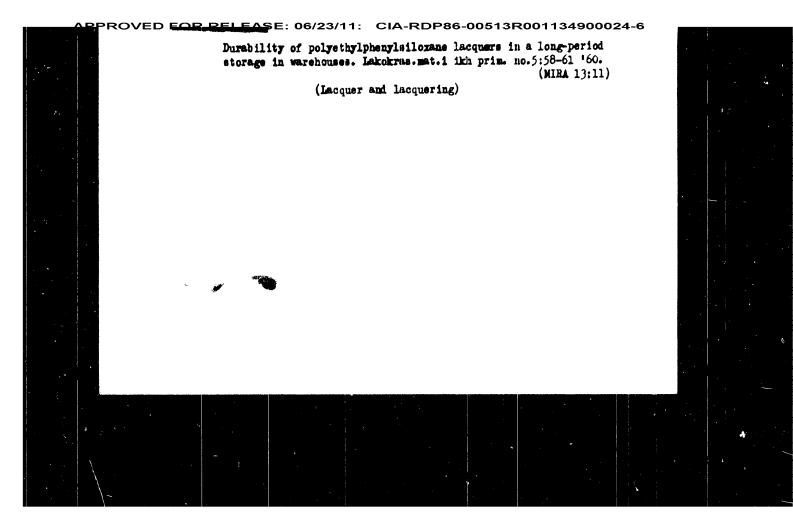
TITLE:

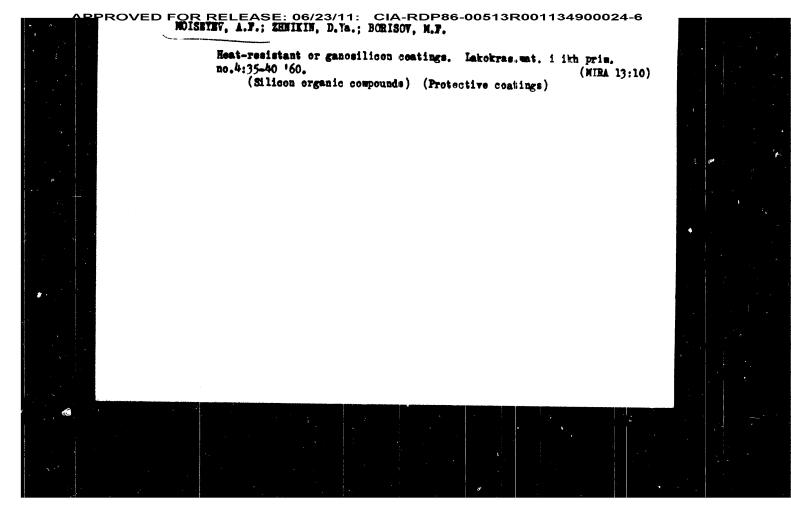
Synthesis of linear polyurethanes without solvents Communication I. Synthesis of linear polyurethanes in the

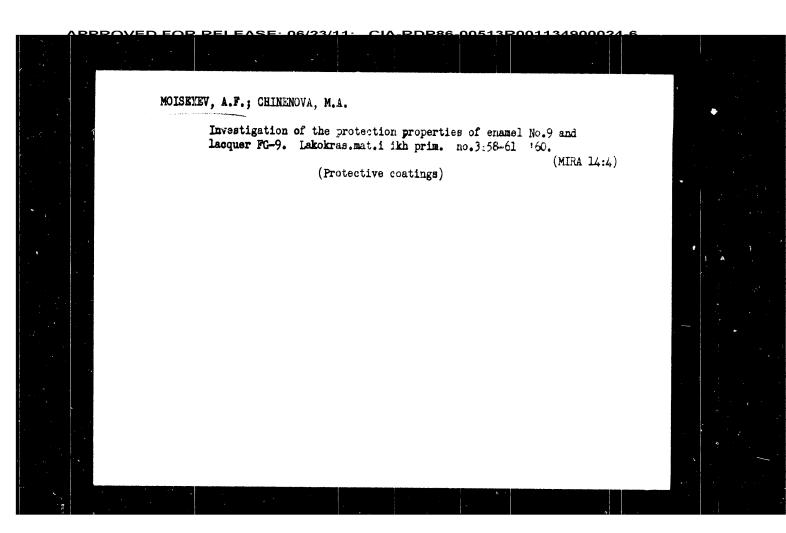
Plasticheskiye massy, no. 6, 1961, 10-11 PERIODICAL:

TEXT: Results are reported according to which the synthesis of polyurethanes in the melt has considerable advantages over the synthesis in inert solvents. The reaction of 1,4-butanediol with hexamethylene dissocyanate was examined. Mixing of equimolar amounts of the two reagents causes the destruction of the polymer due to a great increase in temperature. In a nitrogen atmosphere, colorless polymers were obtained at 23000 with a molecular weight varying from 15,000 to 25,000. In a nitrogen atmosphere free from oxygen, hexamethylene diisocyanate was therefore added dropwise to butanediol heated at 80-90°C. After addition of 90% of the disocyanate the strongly viscous mass was heated at 200-210°C for better stirring, and then, after adding the remaining diisocyanate, the above Card 1/4

APPROVED FOR RELEASE: 06/23/11: CIA-RDP86-00513R001134900024-6







PROVED FOR RELEASE: 06/23/11: CIA-RDP86-00513R001134900024-6 On the Type of Viscosity Change in S/191/60/000/002/006/012 Solutions of the Polyethyl-phenyl-siloxane B027/B058 Resin During Aging in the presence of lead-manganese linoleate than in solutions from pure resin. During ultraviolet irradiation the viscosity change proceeded analogously, but quicker. The quicker viscosity reduction at an addition of lead-manganese linoleate can be explained with the sorption of the molecules of this salt by the dissolved polysiloxane chain molecules. The reciprocal molecular action is thus reduced, the structural bonds between the chain-polymer molecules are destroyed and the viscosity of the solution is reduced. This mechanism was determined by V. A. Kargin and I. V. Mikhaylov during the effect of certain electrolytes on the viscosity of ether cellulose solutions. There are 5 figures and 17 references: 14 Soviet and 3 German.

S/191/60/000/002/006/012 B027/B058

AUTHOR:

Moiseyev, A. F.

TITLE:

On the Type of Viscosity Change in Solutions of the Poly-

ethyl-phenyl-siloxane Resin During Aging

PERIODICAL:

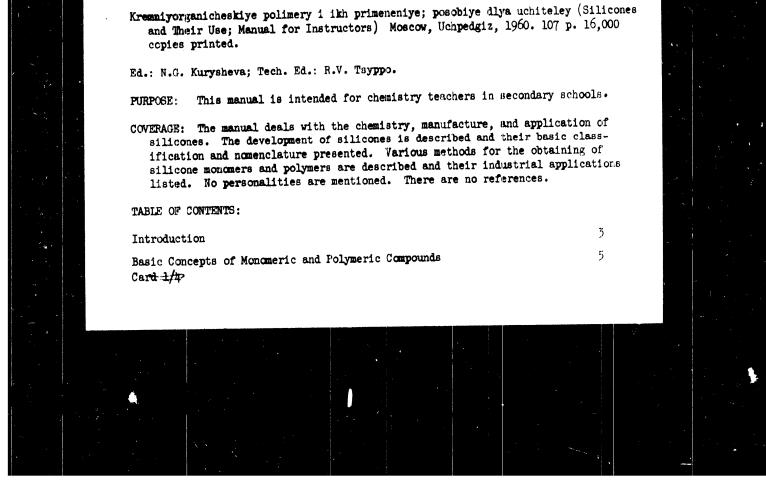
Plasticheskiye massy, 1960, No. 2, pp. 24-26

TEXT: The author studied the viscosity change in solutions of polyethyl-phenyl-siloxane resin in toluene during thermal aging at 60°C and ultraviolet irradiation at 50 to 55°C by the mercury lamp of the type \(\text{PK} - 2 \) (PRK-2). Aging tests were conducted with 43 to 44% solutions of two samples in toluene, i.e. of a pure polyethyl-phenyl-siloxane resin and a resin with an addition of lead-manganese linoleate and siccative 64-6 (64-B). It was established by periodic measurements during thermal aging and subsequent storage of the solutions at normal temperature that the viscosity decreased quickly at the beginning and remained almost stable during further heating. The viscosity of solutions subjected to thermal aging was not re-established at normal temperature during six months. The viscosity reduction in resin solutions of equal concentration is stronger

Card 1/2

ROVED FOR RELEASE: 96/23411년 CLA-RDP86-005/13R001134900024-6 Organosilicon Enamels No. 315 and Γ -4 (G-4). Modification with these resine was carried out by joint condensation of a mixture of the products of joint hydrolysis with polyesters at increased temperature. The heat resistance was tested on pure varnishes and varnishes with pigment admixture (titanium dioxide, chromium oxide, chromium titanate, cadmium red, and aluminum powder) on a steel surface; the protective action and the physicomechanical properties were estimated, which render their practical application possible. The results obtained with resins No. 315 and G-4, as well as with epoxy resin 9-40 (E-40), showed a low heat resistance of the corresponding varnishes at 300°C. Varnishes No. 1 and 2 were much more resistant with an admixture of cadmium red, titanium dioxide, and chromium oxide than without a pigment, both in the pure state and modified by BMK-5 or NI-150. The resulting enamels endure a temperature of 300°C for more than 300 h. After 300 h of heating at 300°C, the average weight loss of the coat is 5.7% in varnishes No. 1 and 2 in the pure state, and 15-22% in varnishes medified with BMK-5 and NI-150. Varnish No. 2 with aluminum powder as a coat endures a temperature of 400 C for more than 100 h, and shows a weight loss of 20.4%. There are 2 tables and 9 references: 4 Soviet, 2 French, and 2 US. Card 2/2

PROVED FOR RELEASE: 06/23/11: CIA-RDP86-00513R001134900024-6 Moisevev. A. F., Maklashina, T. S., Makarova, L. V., Zhinkin, D. Ya., Konstantinova, N. G. AUTHORS: Thermal Stability of Some Protective Organosilicon Enamels TITLE: PERIODICAL: Plasticheskiye massy, 1960, No. 1, pp. 36-38 TEXT: The authors report on their studies of the heat resistance of protective organosilicon enamels which were in operation at 300 and 400°C for a prolonged period. Two types of polymethyl phenyl siloxane resins were used as binding agents for these enamels; the resins were used in the pure state and modified by organic polymers. The R/Si ratio was 1.7, in resin I, and 1.5 in resin II. Toluene solutions of resin I are named No. 1, of resin II, No. 2. To modify the binding agent, the authors used the following substances: 1) Polyacrylate of the type 5MK-5 (BMK-5) which was added both by the mixing of solutions and on heating. It was previously dissolved in solvent No. 648 (FOCT 4006-48, GOST 4006-48). 2) High-viscous ethyl cellulose HM-150 (NI-150) was added in solution. 3) Polyester resins Card 1/2



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Organo-silicic Resins and Varnishes and Their Application

SOV/64-59-2-4/23

of polyethylphenylsiloxane EF-3 and EF-5, polymethylphenyl-siloxane K-40, K-41, K-43, K-44, K-47 and K-48 (Ref 31) are the mostfrequently used. Among a large number of possibilities of combination and application of (OR) for corrosion-proof coatings the heat-resistant enamel varnish Nr 9 and the polydimethyl-phenylsiloxane enamel varnishes PRKE-13 and PRKE-15 are preferably used in Soviet industries (Ref 31). Plastics om (OR) basis are largely used in the electrical industry, as well as for the production of foam plastics. For their production polymethylphenylsiloxane resins K-40 and K-47 as well as modified (OR) with phenol formaldehyde-, epoxy- and polyuretrane ester resins are the most suitable. There are 73 references, 46 of which are Soviet.

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of Technical Sciences

TITLE:

Organo-silicic Resins and Varnishes and Their Application (Kremniyorganicheskiye smoly i laki i ikh primeneniye)

PERIODICAL:

Khimicheskaya promyshlennost', 1959, Nr 2, pp 106-111 (USSR)

ABSTRACT:

In the USSR organo-silicic resins (OR) are largely used in the production of heat-resistant varnishes. At present, there exist several types of (OR), but only polymethylsiloxane-, polymethylsiloxane-, polymethyl-phenylsiloxane-, and polyethylphenylsiloxane resins are being used in pure state and changed with organic polymers. Temperature resistance, hydrophobic properties as well as chemical and oxidation resistance are the most important properties of the varnish- and resin films produced on the basis of (OR). Organo-silicic resins and varnishes exhibit good dielectric properties depending only little on temperature and current frequency. For this reason they are used for electrical insulation. In the USSR the electrical insulation varnishes

Card 1/2